

1 JC05 Rec'd PCT/PTO 18 MAR 2005

DESCRIPTION

NOVEL URACIL DERIVATIVES AND MEDICINAL USE THEREOF

TECHNICAL FIELD

The present invention relates to novel uracil derivatives or pharmaceutically acceptable salts thereof and pharmaceutical compositions for the treatment of allergic diseases and pruritus which contain any of the uracil derivatives or pharmaceutically acceptable salts thereof as an active ingredient.

BACKGROUND ART

10 Allergic reactions capable of causing allergic diseases are mainly classified into type I to type IV allergic reactions. It is said that type IV allergic reactions play a large part in atopic dermatitis, contact dermatitis, chronic bronchial
15 asthma, psoriasis, graft-versus-host disease and the like. The effectiveness of antihistamines and chemical mediator release inhibitors against these diseases is limited, and these diseases are treated by the use of steroids. In addition, immunosuppressive agents such
20 as cyclosporin and tacrolimus are effective in suppressing rejection after organ transplantation or treating graft-versus-host disease. The employment of these drugs for the treatment of dermatitis is expanded

and tacrolimus has already been clinically used for this treatment (Lancet, 339, 1120 (1992) and J. Invest. Dermatol., 98, 851 (1992)).

However, as to the steroids; their adverse
5 effects such as infectious diseases, adrenal atrophy, osteoporosis, diabetes, failure to thrive of children, and the like are problems. Also in the case of the immunosuppressive agents such as cyclosporin and tacrolimus, the exhibition of adverse effects due to
10 their immunosuppressive effect, such as infectious diseases, diabetes and the like is feared. Severe itch is known as one of the characteristics of the morbidity of atopic dermatitis and treatment thereof is clinically very important. However, the effectiveness
15 of existing antihistamines, chemical mediator release inhibitors, steroids and immunosuppressive agents against the itch due to atopic dermatitis is low.

The present applicant has previously proposed uracil derivatives (see JP-A-8-109171) and hydroquinone
20 derivatives (see Japanese Patent No. 3093170), which suppress type IV allergic reactions. There is still desired the development of a more effective therapeutic agent for allergic diseases, in particular, a therapeutic agent which permits suppression of diseases
25 in which a type IV allergic reaction participates or itch accompanying these diseases.

DISCLOSURE OF THE INVENTION

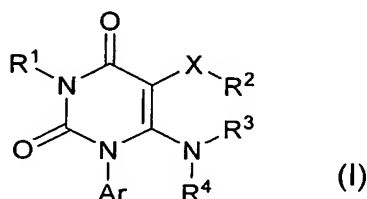
In view of such a situation, an object of the present invention is to provide a novel compound which permits not only treatment of various allergic diseases, in particular, diseases in which a type IV allergic reaction participates, but also suppression of itch accompanying these diseases or suppression of nonallergic itch; and a pharmaceutical composition for the treatment of allergic diseases containing said compound as an active ingredient.

10 In recognition of such a background, in order to develop a therapeutic agent that is more effective against various allergic diseases, in particular, diseases in which a type IV allergic reaction participates, and that can suppress itch accompanying these diseases or nonallergic itch, the present
15 inventors earnestly investigated an aryl group bonded to the 1-position of the uracil skeleton of the uracil derivative disclosed in JP-A-8-109171 and a substituent bonded to the 5-position of this uracil skeleton, and
20 consequently found that the uracil derivative described below in which the aforesaid aryl group is a phenyl group substituted by alkyl groups of 1 to 6 carbon atoms at the o- and m-positions, a substituted or unsubstituted heteroaryl group or a bicyclic aromatic
25 group, and which has a hydroquinone structure with antioxidant action or an structure analogous thereto, which is bonded to the 5-position by a suitable connecting group, suppresses type IV allergic reactions

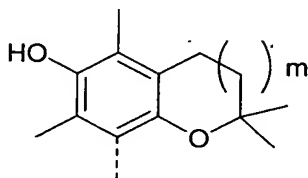
remarkably and also can suppress itch, whereby the present invention has been accomplished.

That is, the present invention includes the following aspects of the invention.

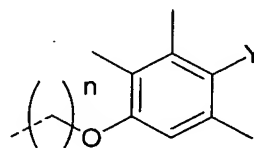
- 5 (1) A uracil derivative represented by the general formula (I):



wherein X represents a group selected from NHCO, NHCH₂, CO, CONH and CH₂NH; R¹ represents a hydrogen atom or a substituted or unsubstituted alkyl group of 1 to 6
 10 carbon atoms; R² represents the general formula (II) or (III):



(II)



(III)

wherein m is 0 or 1, n is an integer of 1 to 3, Y is OH or NH₂, and each dotted line indicates a bonding position, provided that when R² represents the general
 15 formula (III), X represents NHCO or NHCH₂; R³ and R⁴ independently represent a hydrogen atom or an alkyl group of 1 to 6 carbon atoms; and Ar represents a

phenyl group substituted by alkyl groups of 1 to 6 carbon atoms at the o- and m-positions, a substituted or substituted heteroaryl group or a bicyclic aromatic group,

5 or a pharmaceutically acceptable salt thereof.

(2) A uracil derivative or a pharmaceutically acceptable salt thereof according to the above item (1), wherein R^2 represents the general formula (II) in the general formula (I).

10 (3) A uracil derivative or a pharmaceutically acceptable salt thereof according to the above item (1), wherein R^2 represents the general formula (III) in the general formula (I).

(4) A uracil derivative or a pharmaceutically acceptable salt thereof according to the above item (1) 15 or (2), wherein Ar represents a phenyl group substituted by alkyl groups of 1 to 6 carbon atoms at the o- and m-positions in the general formula (I).

(5) A uracil derivative or a pharmaceutically acceptable salt thereof according to the above item (1) 20 or (2), wherein Ar represents a substituted or unsubstituted heteroaryl group in the general formula (I).

(6) A uracil derivative or a pharmaceutically acceptable salt thereof according to the above item (1) 25 or (2), wherein Ar represents a bicyclic aromatic group in the general formula (I).

(7) A uracil derivative or a pharmaceutically

acceptable salt thereof according to any one of the above items (1) to (6), wherein X represents NHCO in the general formula (I).

(8) A pharmaceutical composition for the treatment of allergic diseases comprising a uracil derivative or a pharmaceutically acceptable salt thereof according to any one of the above items (1) to (7) as an active ingredient.

(9) A pharmaceutical composition for the treatment of pruritus comprising a uracil derivative or a pharmaceutically acceptable salt thereof according to any one of the above items (1) to (7) as an active ingredient.

BEST MODE FOR CARRYING OUT THE INVENTION

15 In the present specification, the term "alkyl group of 1 to 6 carbon atoms" means a linear or branched alkyl group of 1 to 6 carbon atoms. Specific examples thereof are methyl group, ethyl group, propyl group (1-propyl group), isopropyl group (2-propyl group), butyl group (1-butyl group), sec-butyl group (2-butyl group), isobutyl group (2-methyl-1-propyl group), t-butyl group (2-methyl-2-propyl group), pentyl group (1-pentyl group) and hexyl group (1-hexyl group). Of these, preferable examples thereof are alkyl groups
20 of 1 to 4 carbon atoms.
25

In the present specification, the term "alkoxy group of 1 to 6 carbon atoms" means a linear or

branched alkoxy group of 1 to 6 carbon atoms. Specific examples thereof are methoxy group, ethoxy group, propoxy group, 1-methylethoxy group, butoxy group, 1-methylpropoxy group, 2-methylpropoxy group, 1,1-
5 dimethylethoxy group, pentyloxy group and hexyloxy group. Of these, preferable examples thereof are alkoxy groups of 1 to 4 carbon atoms.

In the present specification, the halogen atom includes fluorine atom, chlorine atom, bromine
10 atom and iodine atom.

The compound of the present invention is explained below in further detail. When R^2 represents the formula (II), X is a connecting group selected from NHCO, NHCH₂, CO, CONH and CH₂NH and is preferably NHCO.
15 When R^2 represents the formula (III), X is NHCO or NHCH₂.

Preferable examples of the alkyl group of 1 to 6 carbon atoms represented by R^1 are methyl group, ethyl group, propyl group (1-propyl group), isopropyl
20 group (2-propyl group), butyl group (1-butyl group), sec-butyl group (2-butyl group), isobutyl group (2-methyl-1-propyl group) and t-butyl group (2-methyl-2-propyl group).

When the alkyl group of 1 to 6 carbon atoms
25 represented by R^1 is a substituted alkyl group, its substituent(s) includes halogen atoms, hydroxyl group and alkoxy groups of 1 to 6 carbon atoms. The alkyl group may have 1 to 3 substituents which may be the

same or different. Specific examples of the substituents are hydroxyl group, methoxy group, ethoxy group and fluorine atom.

Preferable examples of R^1 are hydrogen atom, methyl group, ethyl group, propyl group (1-propyl group), butyl group (1-butyl group), 2-methoxyethyl group, 2-ethoxyethyl group, 3-methoxypropyl group, 2-hydroxyethyl group, 3-hydroxypropyl group, 4-hydroxybutyl group and 2,2,2-trifluoroethyl group. Of these, hydrogen atom, methyl group, ethyl group, propyl group, butyl group, 2-methoxyethyl group, 2-hydroxyethyl group and 3-hydroxypropyl group are especially preferable.

Preferable examples of each of R^3 and R^4 are hydrogen atom, methyl group, ethyl group and propyl group (1-propyl group). Of these, hydrogen atom and methyl group are especially preferable.

In the phenyl group substituted by alkyl groups of 1 to 6 carbon atoms at the o- and m-positions for Ar, the alkyl groups may be the same or different. Preferable examples of the phenyl group are 2,3-dimethylphenyl group, 2,3-diethylphenyl group, 2,3-dipropylphenyl group, 2,3-dibutylphenyl group, 2-ethyl-3-methylphenyl group, 2-methyl-3-ethylphenyl group and 2-methyl-3-propylphenyl group. Of these, 2,3-dimethylphenyl group is preferable.

As the heteroaryl group for Ar, 5- or 6-membered heteroaryl groups containing one or two

nitrogen atoms are exemplified. Specific examples thereof are 2-pyrrolyl group, 3-pyrrolyl group, 2-pyridyl group, 3-pyridyl group, 4-pyridyl group, pyrimidinyl group and pyrazinyl group. Of these, 2-
5 pyridyl group, 3-pyridyl group and 4-pyridyl group are preferable.

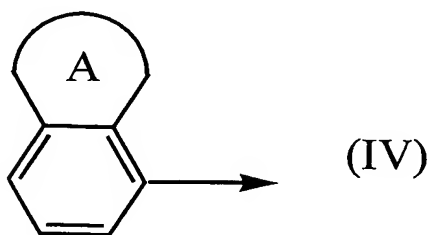
When the heteroaryl group is a substituted group, its substituent(s) includes alkyl groups of 1 to 6 carbon atoms, halogen atoms and alkoxy groups of 1 to
10 6 carbon atoms. Preferable examples of the aforesaid halogen atoms are fluorine atom and chlorine atom. Preferable examples of the aforesaid alkoxy group are methoxy group, ethoxy group, propoxy group, 1-methylethoxy group, butoxy group, 1-methylpropoxy
15 group, 2-methylpropoxy group and 1,1-dimethylethoxy group.

Specific examples of the substituted heteroaryl group are 6-methyl-3-pyridyl group, 6-methoxy-3-pyridyl group, 6-chloro-3-pyridyl group, 2-
20 methyl-3-pyridyl group, 2-methoxy-3-pyridyl group, 2-chloro-3-pyridyl group, 2,6-dimethyl-3-pyridyl group, 2,6-dimethoxyl-3-pyridyl group, 2,6-dichloro-3-pyridyl group and 2,3-dimethyl-4-pyridyl group. Of these, 2-methyl-3-pyridyl group is preferable.

25 As the bicyclic aromatic group, there are exemplified 9- or 10-membered bicyclic aromatic groups that may contain 1 to 3 heteroatoms selected from nitrogen atoms, oxygen atoms and sulfur atoms so that

the number of nitrogen atoms may be 0 to 3, the number of oxygen atoms 0 to 2 and the number of sulfur atoms 0 or 1.

The bicyclic aromatic group preferably refers
5 to a group represented by the formula (IV):



wherein A is a 5- or 6-membered saturated or
unsaturated carbocycle or heterocycle containing 0 to 2
heteroatoms selected from nitrogen atoms, oxygen atoms
and sulfur atoms so that the number of nitrogen atoms
10 may be 0 to 2, the number of oxygen atoms 0 to 2 and
the number of sulfur atoms 0 to 2.

Specific examples of the bicyclic aromatic
group are 1-naphthyl group, 5,6,7,8-tetrahydro-1-
naphthyl group, 2,3-methylenedioxyphenyl group, 2,3-
15 ethylenedioxyphenyl group, 4-indanyl group, 5-quinolyl
group, 5-isoquinolyl group, 5-quinazolinyl group, 5-
quinoxalinyl group, 4-benzofuranyl group, 4-
benzothienyl group, 4-indazolyl group, 4-benzimidazolyl
group and 4-benzothiazolyl group. Of these, 1-naphthyl
20 group, 5,6,7,8-tetrahydro-1-naphthyl group, 4-indanyl
group and 2,3-methylenedioxyphenyl group are
preferable.

Specific preferable examples of the uracil derivative of the present invention are listed in Table 1 to Table 18.

In the following tables, R^1 , R^2 , X, Ar, m, n
5 and Y are as defined in the general formula (I). In this case, each of R^3 and R^4 in the general formula (I) is hydrogen.

Table 1

Compound	R ¹	R ²	X	Ar	m or n	Y
1	Me	(II)	NHCO	1-naphthyl	0	-
2	Me	(II)	NHCO	1-naphthyl	1	-
3	Me	(III)	NHCO	1-naphthyl	1	OH
4	Me	(III)	NHCO	1-naphthyl	1	NH ₂
5	Me	(III)	NHCO	1-naphthyl	2	OH
6	Me	(III)	NHCO	1-naphthyl	2	NH ₂
7	Me	(III)	NHCO	1-naphthyl	3	OH
8	Me	(III)	NHCO	1-naphthyl	3	NH ₂
9	H	(II)	NHCO	1-naphthyl	0	-
10	H	(II)	NHCO	1-naphthyl	1	-
11	H	(III)	NHCO	1-naphthyl	1	NH ₂
12	Et	(III)	NHCO	1-naphthyl	0	NH ₂
13	Pr	(III)	NHCO	1-naphthyl	0	NH ₂
14	Bu	(III)	NHCO	1-naphthyl	0	NH ₂
15	Me	(II)	NHCH ₂	1-naphthyl	0	-
16	Me	(II)	CONH	1-naphthyl	0	-
17	Me	(II)	CO	1-naphthyl	0	-
19	Et	(II)	NHCO	1-naphthyl	0	-
20	Pr	(II)	NHCO	1-naphthyl	0	-
21	Bu	(II)	NHCO	1-naphthyl	0	-
22	Pr	(III)	NHCO	1-naphthyl	1	OH
23	Pr	(III)	NHCO	1-naphthyl	0	NH ₂
24	Pr	(III)	NHCO	1-naphthyl	1	NH ₂
25	Pr	(III)	NHCO	1-naphthyl	0	OH
27	Pr	(III)	NHCO	1-naphthyl	1	NH ₂
28	Pr	(II)	CONH	1-naphthyl	0	-
29	Pr	(II)	NHCH ₂	1-naphthyl	0	-
30	Pr	(II)	CO	1-naphthyl	0	-

Table 2

Compound	R ¹	R ²	X	Ar	m or n	Y
31	H	(II)	NHCO	4-indanyl	0	-
32	H	(II)	NHCO	4-indanyl	1	-
33	H	(III)	NHCO	4-indanyl	1	OH
34	H	(III)	NHCO	4-indanyl	1	NH ₂
35	H	(III)	NHCO	4-indanyl	2	OH
36	H	(III)	NHCO	4-indanyl	2	NH ₂
37	H	(III)	NHCO	4-indanyl	3	OH
38	H	(III)	NHCO	4-indanyl	3	NH ₂
39	H	(II)	NHCH ₂	4-indanyl	0	-
40	H	(II)	NHCH ₂	4-indanyl	1	-
41	H	(III)	NHCH ₂	4-indanyl	1	OH
42	H	(III)	NHCH ₂	4-indanyl	1	NH ₂
43	H	(III)	NHCH ₂	4-indanyl	2	OH
44	H	(III)	NHCH ₂	4-indanyl	2	NH ₂
45	H	(III)	NHCH ₂	4-indanyl	3	OH
46	H	(III)	NHCH ₂	4-indanyl	3	NH ₂
47	H	(II)	CONH	4-indanyl	0	-
48	H	(II)	CONH	4-indanyl	1	-
49	H	(II)	CH ₂ NH	4-indanyl	0	-
50	H	(II)	CH ₂ NH	4-indanyl	1	-
51	H	(II)	CO	4-indanyl	0	-
52	H	(II)	NHCO	4-indanyl	0	-
53	H	(III)	NHCO	4-indanyl	1	OH
54	H	(III)	NHCO	4-indanyl	1	NH ₂
55	H	(II)	NHCO	4-indanyl	0	-
56	H	(III)	NHCO	4-indanyl	1	NH ₂
57	H	(II)	NHCO	4-indanyl	0	-
58	H	(III)	NHCO	4-indanyl	1	NH ₂
59	H	(II)	NHCO	4-indanyl	0	-
60	H	(III)	NHCO	4-indanyl	1	NH ₂

Table 3

Compound	R ¹	R ²	X	Ar	m or n	Y
61	Me	(II)	NHCO	4-indanyl	0	-
62	Me	(II)	NHCO	4-indanyl	1	-
63	Me	(III)	NHCO	4-indanyl	1	OH
64	Me	(III)	NHCO	4-indanyl	1	NH ₂
65	Me	(III)	NHCO	4-indanyl	2	OH
66	Me	(III)	NHCO	4-indanyl	2	NH ₂
67	Me	(III)	NHCO	4-indanyl	3	OH
68	Me	(III)	NHCO	4-indanyl	3	NH ₂
69	Me	(II)	NHCH ₂	4-indanyl	0	-
70	Me	(II)	NHCH ₂	4-indanyl	1	-
71	Me	(III)	NHCH ₂	4-indanyl	1	OH
72	Me	(III)	NHCH ₂	4-indanyl	1	NH ₂
73	Me	(III)	NHCH ₂	4-indanyl	2	OH
74	Me	(III)	NHCH ₂	4-indanyl	2	NH ₂
75	Me	(III)	NHCH ₂	4-indanyl	3	OH
76	Me	(III)	NHCH ₂	4-indanyl	3	NH ₂
77	Me	(II)	CONH	4-indanyl	0	-
78	Me	(II)	CONH	4-indanyl	1	-
79	Me	(II)	CH ₂ NH	4-indanyl	0	-
80	Me	(II)	CH ₂ NH	4-indanyl	1	-
81	Me	(II)	CO	4-indanyl	0	-
82	Me	(II)	NHCO	4-indanyl	0	-
83	Me	(III)	NHCO	4-indanyl	1	OH
84	Me	(III)	NHCO	4-indanyl	1	NH ₂
85	Me	(II)	NHCO	4-indanyl	0	-
86	Me	(III)	NHCO	4-indanyl	1	NH ₂
87	Me	(II)	NHCO	4-indanyl	0	-
88	Me	(III)	NHCO	4-indanyl	1	NH ₂
89	Me	(II)	NHCO	4-indanyl	0	-
90	Me	(III)	NHCO	4-indanyl	1	NH ₂

Table 4

Compound	R ¹	R ²	X	Ar	m or n	Y
91	Et	(II)	NHCO	4-indanyl	0	-
92	Et	(II)	NHCO	4-indanyl	1	-
93	Et	(III)	NHCO	4-indanyl	1	OH
94	Et	(III)	NHCO	4-indanyl	1	NH ₂
95	Et	(III)	NHCO	4-indanyl	2	OH
96	Et	(III)	NHCO	4-indanyl	2	NH ₂
97	Et	(III)	NHCO	4-indanyl	3	OH
98	Et	(III)	NHCO	4-indanyl	3	NH ₂
99	Et	(II)	NHCH ₂	4-indanyl	0	-
100	Et	(II)	NHCH ₂	4-indanyl	1	-
101	Et	(III)	NHCH ₂	4-indanyl	1	OH
102	Et	(III)	NHCH ₂	4-indanyl	1	NH ₂
103	Et	(III)	NHCH ₂	4-indanyl	2	OH
104	Et	(III)	NHCH ₂	4-indanyl	2	NH ₂
105	Et	(III)	NHCH ₂	4-indanyl	3	OH
106	Et	(III)	NHCH ₂	4-indanyl	3	NH ₂
107	Et	(II)	CONH	4-indanyl	0	-
108	Et	(II)	CONH	4-indanyl	1	-
109	Et	(II)	CH ₂ NH	4-indanyl	0	-
110	Et	(II)	CH ₂ NH	4-indanyl	1	-
111	Et	(II)	CO	4-indanyl	0	-
112	Et	(II)	NHCO	4-indanyl	0	-
113	Et	(III)	NHCO	4-indanyl	1	OH
114	Et	(III)	NHCO	4-indanyl	1	NH ₂
115	Et	(II)	NHCO	4-indanyl	0	-
116	Et	(III)	NHCO	4-indanyl	1	NH ₂
117	Et	(II)	NHCO	4-indanyl	0	-
118	Et	(III)	NHCO	4-indanyl	1	NH ₂
119	Et	(II)	NHCO	4-indanyl	0	-
120	Et	(III)	NHCO	4-indanyl	1	NH ₂

Table 5

Compound	R ¹	R ²	X	Ar	m or n	Y
121	Pr	(II)	NHCO	4-indanyl	0	-
122	Pr	(II)	NHCO	4-indanyl	1	-
123	Pr	(III)	NHCO	4-indanyl	1	OH
124	Pr	(III)	NHCO	4-indanyl	1	NH ₂
125	Pr	(III)	NHCO	4-indanyl	2	OH
126	Pr	(III)	NHCO	4-indanyl	2	NH ₂
127	Pr	(III)	NHCO	4-indanyl	3	OH
128	Pr	(III)	NHCO	4-indanyl	3	NH ₂
129	Pr	(II)	NHCH ₂	4-indanyl	0	-
130	Pr	(II)	NHCH ₂	4-indanyl	1	-
131	Pr	(III)	NHCH ₂	4-indanyl	1	OH
132	Pr	(III)	NHCH ₂	4-indanyl	1	NH ₂
133	Pr	(III)	NHCH ₂	4-indanyl	2	OH
134	Pr	(III)	NHCH ₂	4-indanyl	2	NH ₂
135	Pr	(III)	NHCH ₂	4-indanyl	3	OH
136	Pr	(III)	NHCH ₂	4-indanyl	3	NH ₂
137	Pr	(II)	CONH	4-indanyl	0	-
138	Pr	(II)	CONH	4-indanyl	1	-
139	Pr	(II)	CH ₂ NH	4-indanyl	0	-
140	Pr	(II)	CH ₂ NH	4-indanyl	1	-
141	Pr	(II)	CO	4-indanyl	0	-
122	Pr	(II)	NHCO	4-indanyl	0	-
143	Pr	(III)	NHCO	4-indanyl	1	OH
144	Pr	(III)	NHCO	4-indanyl	1	NH ₂
145	Pr	(II)	NHCO	4-indanyl	0	-
146	Pr	(III)	NHCO	4-indanyl	1	NH ₂
147	Pr	(II)	NHCO	4-indanyl	0	-
148	Pr	(III)	NHCO	4-indanyl	1	NH ₂
149	Pr	(II)	NHCO	4-indanyl	0	-
150	Pr	(III)	NHCO	4-indanyl	1	NH ₂

Table 6

Compound	R ¹	R ²	X	Ar	m or n	Y
151	H	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
152	H	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	-
153	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	OH
154	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
155	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	2	OH
156	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	2	NH ₂
157	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	3	OH
158	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	3	NH ₂
159	H	(II)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	0	-
160	H	(II)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	-
161	H	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	OH
162	H	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
163	H	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	2	OH
164	H	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	2	NH ₂
165	H	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	3	OH

- Contin'd -

- Contin'd -

166	H	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	3	NH ₂
167	H	(II)	CONH	5,6,7,8-tetrahydro-1-naphthyl	0	-
168	H	(II)	CONH	5,6,7,8-tetrahydro-1-naphthyl	1	-
169	H	(II)	CH ₂ NH	5,6,7,8-tetrahydro-1-naphthyl	0	-
170	H	(II)	CH ₂ NH	5,6,7,8-tetrahydro-1-naphthyl	1	-
171	H	(II)	CO	5,6,7,8-tetrahydro-1-naphthyl	0	-
172	H	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
173	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	OH
174	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
175	H	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
176	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
177	H	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
178	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
179	H	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
180	H	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂

Table 7

Compound	R ¹	R ²	X	Ar	m or n	Y
181	Me	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
182	Me	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	-
183	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	OH
184	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
185	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	2	OH
186	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	2	NH ₂
187	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	3	OH
188	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	3	NH ₂
189	Me	(II)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	0	-
190	Me	(II)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	-
191	Me	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	OH
192	Me	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
193	Me	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	2	OH
194	Me	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	2	NH ₂
195	Me	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	3	OH

- Contin'd -

- Contin'd -

196	Me	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	3	NH ₂
197	Me	(II)	CONH	5,6,7,8-tetrahydro-1-naphthyl	0	-
198	Me	(II)	CONH	5,6,7,8-tetrahydro-1-naphthyl	1	-
199	Me	(II)	CH ₂ NH	5,6,7,8-tetrahydro-1-naphthyl	0	-
200	Me	(II)	CH ₂ NH	5,6,7,8-tetrahydro-1-naphthyl	1	-
201	Me	(II)	CO	5,6,7,8-tetrahydro-1-naphthyl	0	-
202	Me	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
203	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	OH
204	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
205	Me	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
206	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
207	Me	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
208	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
209	Me	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
210	Me	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂

Table 8

Compound	R ¹	R ²	X	Ar	m or n	Y
211	Et	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
212	Et	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	-
213	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	OH
214	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
215	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	2	OH
216	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	2	NH ₂
217	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	3	OH
218	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	3	NH ₂
219	Et	(II)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	0	-
220	Et	(II)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	-
221	Et	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	OH
222	Et	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
223	Et	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	2	OH
224	Et	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	2	NH ₂
225	Et	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	3	OH

- Contin'd -

- Contin'd -

226	Et	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	3	NH ₂
227	Et	(II)	CONH	5,6,7,8-tetrahydro-1-naphthyl	0	-
228	Et	(II)	CONH	5,6,7,8-tetrahydro-1-naphthyl	1	-
229	Et	(II)	CH ₂ NH	5,6,7,8-tetrahydro-1-naphthyl	0	-
230	Et	(II)	CH ₂ NH	5,6,7,8-tetrahydro-1-naphthyl	1	-
231	Et	(II)	CO	5,6,7,8-tetrahydro-1-naphthyl	0	-
232	Et	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
233	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	OH
234	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
235	Et	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
236	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
237	Et	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
238	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
239	Et	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
240	Et	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂

Table 9

Compound	R ¹	R ²	X	Ar	m or n	Y
241	Pr	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
242	Pr	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	-
243	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	OH
244	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
245	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	2	OH
246	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	2	NH ₂
247	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	3	OH
248	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	3	NH ₂
249	Pr	(II)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	0	-
250	Pr	(II)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	-
251	Pr	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	OH
252	Pr	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
253	Pr	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	2	OH
254	Pr	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	2	NH ₂
255	Pr	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	3	OH

- Contin'd -

- Contin'd -

256	Pr	(III)	NHCH ₂	5,6,7,8-tetrahydro-1-naphthyl	3	NH ₂
257	Pr	(II)	CONH	5,6,7,8-tetrahydro-1-naphthyl	0	-
258	Pr	(II)	CONH	5,6,7,8-tetrahydro-1-naphthyl	1	-
259	Pr	(II)	CH ₂ NH	5,6,7,8-tetrahydro-1-naphthyl	0	-
260	Pr	(II)	CH ₂ NH	5,6,7,8-tetrahydro-1-naphthyl	1	-
261	Pr	(II)	CO	5,6,7,8-tetrahydro-1-naphthyl	0	-
262	Pr	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
263	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	OH
264	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
265	Pr	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
266	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
267	Pr	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
268	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂
269	Pr	(II)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	0	-
270	Pr	(III)	NHCO	5,6,7,8-tetrahydro-1-naphthyl	1	NH ₂

Table 10

Compound	R ¹	R ²	X	Ar	m or n	Y
271	H	(II)	NHCO	2,3-dimethylphenyl	0	-
272	H	(II)	NHCO	2,3-dimethylphenyl	1	-
273	H	(III)	NHCO	2,3-dimethylphenyl	1	OH
274	H	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
275	H	(III)	NHCO	2,3-dimethylphenyl	2	OH
276	H	(III)	NHCO	2,3-dimethylphenyl	2	NH ₂
277	H	(III)	NHCO	2,3-dimethylphenyl	3	OH
278	H	(III)	NHCO	2,3-dimethylphenyl	3	NH ₂
279	H	(II)	NHCH ₂	2,3-dimethylphenyl	0	-
280	H	(II)	NHCH ₂	2,3-dimethylphenyl	1	-
281	H	(III)	NHCH ₂	2,3-dimethylphenyl	1	OH
282	H	(III)	NHCH ₂	2,3-dimethylphenyl	1	NH ₂
283	H	(III)	NHCH ₂	2,3-dimethylphenyl	2	OH
284	H	(III)	NHCH ₂	2,3-dimethylphenyl	2	NH ₂
285	H	(III)	NHCH ₂	2,3-dimethylphenyl	3	OH
286	H	(III)	NHCH ₂	2,3-dimethylphenyl	3	NH ₂
287	H	(II)	CONH	2,3-dimethylphenyl	0	-
288	H	(II)	CONH	2,3-dimethylphenyl	1	-
289	H	(II)	CH ₂ NH	2,3-dimethylphenyl	0	-
290	H	(II)	CH ₂ NH	2,3-dimethylphenyl	1	-
291	H	(II)	CO	2,3-dimethylphenyl	0	-
292	H	(II)	NHCO	2,3-dimethylphenyl	0	-

- Contin'd -

- Contin'd -

293	H	(III)	NHCO	2,3- dimethylphenyl	1	OH
294	H	(III)	NHCO	2,3- dimethylphenyl	1	NH ₂
295	H	(II)	NHCO	2,3- dimethylphenyl	0	-
296	H	(III)	NHCO	2,3- dimethylphenyl	1	NH ₂
297	H	(II)	NHCO	2,3- dimethylphenyl	0	-
298	H	(III)	NHCO	2,3- dimethylphenyl	1	NH ₂
299	H	(II)	NHCO	2,3- dimethylphenyl	0	-
300	H	(III)	NHCO	2,3- dimethylphenyl	1	NH ₂

Table 11

Compound	R ¹	R ²	X	Ar	m or n	Y
301	Me	(II)	NHCO	2,3-dimethylphenyl	0	-
302	Me	(II)	NHCO	2,3-dimethylphenyl	1	-
303	Me	(III)	NHCO	2,3-dimethylphenyl	1	OH
304	Me	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
305	Me	(III)	NHCO	2,3-dimethylphenyl	2	OH
306	Me	(III)	NHCO	2,3-dimethylphenyl	2	NH ₂
307	Me	(III)	NHCO	2,3-dimethylphenyl	3	OH
308	Me	(III)	NHCO	2,3-dimethylphenyl	3	NH ₂
309	Me	(II)	NHCH ₂	2,3-dimethylphenyl	0	-
310	Me	(II)	NHCH ₂	2,3-dimethylphenyl	1	-
311	Me	(III)	NHCH ₂	2,3-dimethylphenyl	1	OH
312	Me	(III)	NHCH ₂	2,3-dimethylphenyl	1	NH ₂
313	Me	(III)	NHCH ₂	2,3-dimethylphenyl	2	OH
314	Me	(III)	NHCH ₂	2,3-dimethylphenyl	2	NH ₂
315	Me	(III)	NHCH ₂	2,3-dimethylphenyl	3	OH
316	Me	(III)	NHCH ₂	2,3-dimethylphenyl	3	NH ₂
317	Me	(II)	CONH	2,3-dimethylphenyl	0	-
318	Me	(II)	CONH	2,3-dimethylphenyl	1	-
319	Me	(II)	CH ₂ NH	2,3-dimethylphenyl	0	-
320	Me	(II)	CH ₂ NH	2,3-dimethylphenyl	1	-
321	Me	(II)	CO	2,3-dimethylphenyl	0	-
322	Me	(II)	NHCO	2,3-dimethylphenyl	0	-

- Contin'd -

- Contin'd -

323	Me	(III)	NHCO	2,3-dimethylphenyl	1	OH
324	Me	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
325	Me	(II)	NHCO	2,3-dimethylphenyl	0	-
326	Me	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
327	Me	(II)	NHCO	2,3-dimethylphenyl	0	-
328	Me	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
329	Me	(II)	NHCO	2,3-dimethylphenyl	0	-
330	Me	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂

Table 12

Compound	R ¹	R ²	X	Ar	m or n	Y
331	Et	(II)	NHCO	2,3-dimethylphenyl	0	-
332	Et	(II)	NHCO	2,3-dimethylphenyl	1	-
333	Et	(III)	NHCO	2,3-dimethylphenyl	1	OH
334	Et	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
335	Et	(III)	NHCO	2,3-dimethylphenyl	2	OH
336	Et	(III)	NHCO	2,3-dimethylphenyl	2	NH ₂
337	Et	(III)	NHCO	2,3-dimethylphenyl	3	OH
338	Et	(III)	NHCO	2,3-dimethylphenyl	3	NH ₂
339	Et	(II)	NHCH ₂	2,3-dimethylphenyl	0	-
340	Et	(II)	NHCH ₂	2,3-dimethylphenyl	1	-
341	Et	(III)	NHCH ₂	2,3-dimethylphenyl	1	OH
342	Et	(III)	NHCH ₂	2,3-dimethylphenyl	1	NH ₂
343	Et	(III)	NHCH ₂	2,3-dimethylphenyl	2	OH
344	Et	(III)	NHCH ₂	2,3-dimethylphenyl	2	NH ₂
345	Et	(III)	NHCH ₂	2,3-dimethylphenyl	3	OH
346	Et	(III)	NHCH ₂	2,3-dimethylphenyl	3	NH ₂
347	Et	(II)	CONH	2,3-dimethylphenyl	0	-
348	Et	(II)	CONH	2,3-dimethylphenyl	1	-
349	Et	(II)	CH ₂ NH	2,3-dimethylphenyl	0	-

- Contin'd -

- Contin'd -

350	Et	(II)	CH ₂ NH	2,3-dimethylphenyl	1	-
351	Et	(II)	CO	2,3-dimethylphenyl	0	-
352	Et	(II)	NHCO	2,3-dimethylphenyl	0	-
353	Et	(III)	NHCO	2,3-dimethylphenyl	1	OH
354	Et	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
355	Et	(II)	NHCO	2,3-dimethylphenyl	0	-
356	Et	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
357	Et	(II)	NHCO	2,3-dimethylphenyl	0	-
358	Et	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
359	Et	(II)	NHCO	2,3-dimethylphenyl	0	-
360	Et	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂

Table 13

Compound	R ¹	R ²	X	Ar	m or n	Y
361	Pr	(II)	NHCO	2,3-dimethylphenyl	0	-
362	Pr	(II)	NHCO	2,3-dimethylphenyl	1	-
363	Pr	(III)	NHCO	2,3-dimethylphenyl	1	OH
364	Pr	(III)	NHCO	2,3-dimethylphenyl	1	NH ₂
365	Pr	(III)	NHCO	2,3-dimethylphenyl	2	OH
366	Pr	(III)	NHCO	2,3-dimethylphenyl	2	NH ₂
367	Pr	(III)	NHCO	2,3-dimethylphenyl	3	OH
368	Pr	(III)	NHCO	2,3-dimethylphenyl	3	NH ₂
369	Pr	(II)	NHCH ₂	2,3-dimethylphenyl	0	-
370	Pr	(II)	NHCH ₂	2,3-dimethylphenyl	1	-
371	Pr	(III)	NHCH ₂	2,3-dimethylphenyl	1	OH
372	Pr	(III)	NHCH ₂	2,3-dimethylphenyl	1	NH ₂
373	Pr	(III)	NHCH ₂	2,3-dimethylphenyl	2	OH
374	Pr	(III)	NHCH ₂	2,3-dimethylphenyl	2	NH ₂
375	Pr	(III)	NHCH ₂	2,3-dimethylphenyl	3	OH
376	Pr	(III)	NHCH ₂	2,3-dimethylphenyl	3	NH ₂
377	Pr	(II)	CONH	2,3-dimethylphenyl	0	-
378	Pr	(II)	CONH	2,3-dimethylphenyl	1	-
379	Pr	(II)	CH ₂ NH	2,3-dimethylphenyl	0	-
380	Pr	(II)	CH ₂ NH	2,3-dimethylphenyl	1	-
381	Pr	(II)	CO	2,3-dimethylphenyl	0	-
382	Pr	(II)	NHCO	2,3-dimethylphenyl	0	-

- Contin'd -

- Contin'd -

383	Pr	(III)	NHCO	2,3- dimethylphenyl	1	OH
384	Pr	(III)	NHCO	2,3- dimethylphenyl	1	NH ₂
385	Pr	(II)	NHCO	2,3- dimethylphenyl	0	-
386	Pr	(III)	NHCO	2,3- dimethylphenyl	1	NH ₂
387	Pr	(II)	NHCO	2,3- dimethylphenyl	0	-
388	Pr	(III)	NHCO	2,3- dimethylphenyl	1	NH ₂
389	Pr	(II)	NHCO	2,3- dimethylphenyl	0	-
390	Pr	(III)	NHCO	2,3- dimethylphenyl	1	NH ₂

Table 14

Compound	R ¹	R ²	X	Ar	m or n	Y
391	Me	(II)	NHCO	3-Py	0	-
392	Me	(II)	NHCO	3-Py	1	-
393	Me	(III)	NHCO	3-Py	1	OH
394	Me	(III)	NHCO	3-Py	1	NH ₂
395	Me	(III)	NHCO	3-Py	2	OH
396	Me	(III)	NHCO	3-Py	2	NH ₂
397	Me	(III)	NHCO	3-Py	3	OH
398	Me	(III)	NHCO	3-Py	3	NH ₂
399	Et	(II)	NHCO	3-Py	0	-
400	Et	(II)	NHCO	3-Py	1	-
401	Et	(III)	NHCO	3-Py	1	OH
402	Et	(III)	NHCO	3-Py	1	NH ₂
403	Et	(III)	NHCO	3-Py	2	OH
404	Et	(III)	NHCO	3-Py	2	NH ₂
405	Et	(III)	NHCO	3-Py	3	OH
406	Et	(III)	NHCO	3-Py	3	NH ₂
407	Pr	(II)	NHCO	3-Py	0	-
408	Pr	(II)	NHCO	3-Py	1	-
409	Pr	(III)	NHCO	3-Py	1	OH
410	Pr	(III)	NHCO	3-Py	0	NH ₂
411	Pr	(III)	NHCO	3-Py	1	OH
412	Pr	(III)	NHCO	3-Py	0	NH ₂
413	Pr	(III)	NHCO	3-Py	1	NH ₂
414	Pr	(III)	NHCO	3-Py	0	OH
415	Pr	(III)	NHCO	3-Py	1	NH ₂

Table 15

Compound	R ¹	R ²	X	Ar	m or n	Y
416	Me	(II)	NHCO	2-Py	0	-
417	Me	(II)	NHCO	2-Py	1	-
418	Et	(III)	NHCO	2-Py	0	OH
419	Et	(III)	NHCO	2-Py	1	NH ₂
420	Pr	(III)	NHCO	2-Py	0	OH
421	Pr	(III)	NHCO	2-Py	1	NH ₂
422	Me	(III)	NHCO	2-Me-3-Py	0	OH
423	Me	(III)	NHCO	2-Me-3-Py	1	NH ₂
424	Et	(II)	NHCO	2-Me-3-Py	0	-
425	Et	(II)	NHCO	2-Me-3-Py	1	-
426	Pr	(III)	NHCO	2-Me-3-Py	0	OH
427	Pr	(III)	NHCO	2-Me-3-Py	1	NH ₂
428	Me	(III)	NHCO	2-OMe-3-Py	0	OH
429	Me	(III)	NHCO	2-OMe-3-Py	1	NH ₂
430	Et	(III)	NHCO	2-OMe-3-Py	0	OH
431	Et	(III)	NHCO	2-OMe-3-Py	1	NH ₂
432	Pr	(II)	NHCO	2-OMe-3-Py	0	-
433	Pr	(II)	NHCO	2-OMe-3-Py	1	-
434	Me	(III)	NHCO	6-OMe-Py	0	OH
435	Me	(III)	NHCO	6-OMe-Py	1	NH ₂
436	Et	(III)	NHCO	6-OMe-Py	0	OH
437	Et	(III)	NHCO	6-OMe-Py	1	NH ₂
438	Et	(III)	NHCO	6-OMe-Py	0	-
439	Et	(III)	NHCO	6-OMe-Py	1	-

Table 16

Compound	R ¹	R ²	X	Ar	m or n	Y
440	Me	(II)	NHCO	4-Py	0	-
441	Me	(II)	NHCO	4-Py	1	-
442	Et	(III)	NHCO	4-Py	0	OH
443	Et	(III)	NHCO	4-Py	1	NH ₂
444	Pr	(III)	NHCO	4-Py	0	OH
445	Pr	(III)	NHCO	4-Py	1	NH ₂
446	Me	(III)	NHCO	2,6-OMe-3-Py	0	OH
447	Me	(III)	NHCO	2,6-OMe-3-Py	1	NH ₂
448	Et	(II)	NHCO	2,6-OMe-3-Py	0	-
449	Et	(II)	NHCO	2,6-OMe-3-Py	1	-
450	Et	(III)	NHCO	2,6-OMe-3-Py	0	OH
451	Et	(III)	NHCO	2,6-OMe-3-Py	1	NH ₂
452	Me	(III)	NHCO	6-Cl-3-Py	0	OH
453	Me	(III)	NHCO	6-Cl-3-Py	1	NH ₂
454	Et	(III)	NHCO	6-Cl-3-Py	0	OH
455	Et	(III)	NHCO	6-Cl-3-Py	1	NH ₂
456	Et	(II)	NHCO	6-Cl-3-Py	0	-
457	Et	(II)	NHCO	6-Cl-3-Py	1	-
458	Me	(III)	NHCO	2,3-diethylphenyl	0	OH
459	Me	(III)	NHCO	2,3-diethylphenyl	1	NH ₂
460	Et	(III)	NHCO	2,3-diethylphenyl	0	OH
461	Et	(III)	NHCO	2,3-diethylphenyl	1	NH ₂
462	Pr	(III)	NHCO	2,3-diethylphenyl	0	OH
463	Pr	(III)	NHCO	2,3-diethylphenyl	1	NH ₂

Table 17

Compound	R ¹	R ²	X	Ar	m or n	Y
464	(CH ₂) 2OMe	(II)	NHCO	2,3-Me-Ph	0	-
465	(CH ₂) 3OMe	(II)	NHCO	2,3-Me-Ph	0	-
466	(CH ₂) 2OH	(II)	NHCO	2,3-Me-Ph	0	-
467	(CH ₂) 3OH	(II)	NHCO	2,3-Me-Ph	0	-
468	(CH ₂) 4OH	(II)	NHCO	2,3-Me-Ph	0	-
469	(CH ₂) 2OH	(II)	NHCO	2,3-Me-Ph	1	-
470	(CH ₂) 3OH	(II)	NHCO	2,3-Me-Ph	1	-
471	(CH ₂) 4OH	(II)	NHCO	2,3-Me-Ph	1	-
472	(CH ₂) 2OH	(III)	NHCO	2,3-Me-Ph	1	NH ₂
473	(CH ₂) 3OH	(III)	NHCO	2,3-Me-Ph	1	NH ₂
474	(CH ₂) 4OH	(III)	NHCO	2,3-Me-Ph	1	NH ₂
475	(CH ₂) 2OH	(II)	NHCO	4-indanyl	0	-
476	(CH ₂) 3OH	(II)	NHCO	4-indanyl	0	-
477	(CH ₂) 4OH	(II)	NHCO	4-indanyl	0	-
478	(CH ₂) 2OH	(II)	NHCO	5,6,7,8-tetrahydro -1-naphthyl	0	-
479	(CH ₂) 3OH	(II)	NHCO	5,6,7,8-tetrahydro -1-naphthyl	0	-
480	(CH ₂) 4OH	(II)	NHCO	5,6,7,8-tetrahydro -1-naphthyl	0	-

In the following table, R^2 , X, Ar, m, n, R^4 and Y are as defined in the general formula (I). In this case, in the general formula (I), R^1 is a methyl group and R^3 is hydrogen.

Table 18

Compound	R^1	R^2	X	Ar	m or n	Y
481	(III)	NHCO	2,3-dimethyl phenyl	1	Me	NH ₂
482	(III)	NHCO	2,3-dimethyl phenyl	1	Et	NH ₂
483	(III)	NHCO	2,3-dimethyl phenyl	1	Pr	NH ₂
484	(III)	NHCO	2,3-dimethyl phenyl	1	Bu	NH ₂
485	(III)	NHCO	2,3-dimethyl phenyl	1	i-Bu	NH ₂
486	(III)	NHCO	2,3-dimethyl phenyl	1	H	NH ₂
487	(III)	NHCO	1-naphthyl	1	Me	NH ₂
488	(III)	NHCO	1-naphthyl	1	Et	NH ₂
489	(III)	NHCO	1-naphthyl	1	Pr	NH ₂
490	(III)	NHCO	1-naphthyl	1	Bu	NH ₂
491	(III)	NHCO	1-naphthyl	1	i-Bu	NH ₂
492	(III)	NHCO	4-indanyl	1	Me	NH ₂
493	(III)	NHCO	4-indanyl	1	Et	NH ₂
494	(III)	NHCO	4-indanyl	1	Pr	NH ₂
495	(III)	NHCO	4-indanyl	1	Bu	NH ₂

- Contin'd -

496	(III)	NHCO	4-indanyl	1	i-Bu	NH ₂
497	(III)	NHCO	2,3-dimethyl phenyl	1	Me	OH
498	(III)	NHCO	2,3-dimethyl phenyl	1	Et	OH
499	(III)	NHCO	2,3-dimethyl phenyl	1	Pr	OH
500	(III)	NHCO	2,3-dimethyl phenyl	1	Bu	OH
501	(III)	NHCO	2,3-dimethyl phenyl	1	i-Bu	OH
502	(II)	NHCO	1-naphthyl	0	Me	-
503	(III)	NHCO	4-indanyl	1	Et	OH
504	(III)	NHCO	4-indanyl	1	Pr	OH
505	(III)	NHCO	4-indanyl	1	Bu	OH
506	(II)	NHCO	2,3-dimethyl phenyl	0	Me	-

The uracil derivative represented by the general formula (I) may be synthesized by combining a 1-substituted-2,4(1H,3H)-3-(substituted or unsubstituted)-pyrimidinedione derivative, an intermediate for the synthesis with a corresponding antioxidant unit under suitable reaction conditions. Specifically, a compound of the formula (I) in which X = NHCO may be synthesized by amidation reaction of a 5,6-diamino-3-(substituted or unsubstituted)-1-substituted-2,4(1H,3H)-pyrimidinedione derivative with

a corresponding carboxylic acid. For example, an acid halide converted from the carboxylic acid may be used in the amidation reaction. It is also possible to react the carboxylic acid with an activated reagent
 5 such as a carbodiimide or a phosphoryl halide and use the reaction product in the amidation reaction. A compound of the formula (I) in which $X = \text{NHCH}_2$ may be synthesized by synthesizing a corresponding compound of the formula (I) in which $X = \text{NHCO}$ by the process
 10 described above, and reducing this compound with diborane.

A compound of the formula (I) in which $X = \text{CONH}$ and a compound of the formula (I) in which $X = \text{CO}$ may be synthesized by a process based on the method of
 15 Bernier et al. (Bull. Soc. Chim. Fr., 1976, 616). That is, the compound in which $X = \text{CONH}$ may be synthesized by directly reacting a 6-amino-3-(substituted or unsubstituted)-1-substituted-2,4(1H,3H)-pyrimidinedione derivative with an isocyanate. The compound in which X
 20 $= \text{CO}$ may be synthesized by directly reacting a 6-amino-3-(substituted or unsubstituted)-1-substituted-2,4(1H,3H)-pyrimidinedione with an acid halide. A compound of the formula (I) in which $X = \text{CH}_2\text{NH}$ may be synthesized by synthesizing a corresponding compound of
 25 the formula (I) in which $X = \text{CONH}$ by the method described above and reducing this compound with diborane.

3-(Substituted or unsubstituted)-1-

substituted-5,6-diamino-2,4(1H,3H)-pyrimidinedione, an intermediate for the synthesis having a 2,4(1H,3H)-pyrimidinedione ring, may be synthesized, for example, by the same method as described in JP-A-8-109171 and
5 Japanese Patent No. 3093170.

A 1,3-substituted-2,4(1H,3H)-pyrimidinedione in which R^1 is a lower alkyl group may be obtained by converting the amino group at the 6-position to a hydroxyl group with hydrochloric acid, replacing the
10 hydroxyl group with a chloro group, for example, by the method of Senda et al. (Chem. Pharm. Bull., 1974, 22, 189), and then reacting the resulting compound with any of various alkylamines. Thereafter, an amino group may be introduced into the 5-position by the same method as
15 described above.

When R^2 represents the general formula (II) and X is NHCO, a carboxylic acid used as a starting material may be synthesized, for example, by the similar method of Lars et al. (Tetrahedron, 1970, 26,
20 879) ($m = 0$) or the process disclosed in JP-A-7-215959 ($m = 1$). When R^2 represents the general formula (II) and X is CONH, an isocyanate used as a starting material may be synthesized by reacting the carboxylic acid obtained by the above with diphenylphosphoryl
25 azide.

When R^2 represents the general formula (III) and X is NHCO, a starting material may be synthesized by protecting the phenolic hydroxyl group or amino

group of commercial 2,3,5-trimethyl-1,4-hydroquinone and 2,3,5-trimethyl-4-hydroxyaniline (Journal of American Chemical Society, 1939, 61, 765), respectively, by conventional methods; reacting them
5 with a haloalkanoate, and then hydrolyzing the reaction product.

As pharmaceutically acceptable salts of derivatives having a functional group capable of forming the salt, among the uracil derivatives
10 represented by the general formula (I), there are exemplified hydrochloride, sulfate, acetate, succinate, sodium salt, potassium salt, calcium salt and ammonium salt. These salts may be obtained by mixing the uracil derivative with an acid or a base, followed by
15 purification by a conventional method such as recrystallization.

The present invention includes hydrates and solvates (e.g. ethanol solvates) of the uracil derivatives represented by the general formula (I) or
20 pharmaceutically acceptable salts thereof. In addition, the present invention includes all tautomers of the uracil derivatives represented by the general formula (I), all stereoisomers (e.g. optical isomers) which the uracil derivatives have, and all crystal
25 forms of the uracil derivatives.

The therapeutic agent for allergic diseases and pruritus of the present invention may be used in any of dosage forms such as oral preparations (e.g.

tablets, capsules and powders), injections, external preparations and the like. For example, the uracil derivative or pharmaceutically acceptable salt thereof of the present invention may be formulated into an ointment by mixing with an ointment base such as vaseline. The uracil derivative or pharmaceutically acceptable salt thereof of the present invention may also be formulated into tablets by mixing with conventional additives such as excipients (e.g. lactose and starch), lubricants (e.g. magnesium stearate and talc) and the like.

The dose of the therapeutic agent for allergic diseases and pruritus of the present invention is properly determined depending on the sex, age and body weight of a patient, a disease to be treated, the symptom of the patient, and the like. For example, an ointment containing 0.01 to 10% of the active ingredient may be applied to an affected part once to several times per day in the case of a skin disease such as atopic dermatitis, contact dermatitis, psoriasis, pruritus in dialysis, or the like. When used in the form of an oral preparation such as tablets, capsules, powder or the like, the therapeutic agent of the present invention may be administered in a dose of 0.01 to 100 mg/kg per day in one portion or several portions.

EXAMPLES

The present invention is concretely illustrated with the following examples, which should not be construed as limiting the scope of the invention.

5 Reference Example 1: 6-amino-3-methyl-1-(1-naphthyl)uracil

A 40% aqueous methylamine solution (3.82 mL) was added to a solution of 1-naphthyl isocyanate (5.01 g, 29.6 mmol) in acetonitrile (100 mL) under ice-
10 cooling. The temperature was raised to room temperature and the reaction was carried out for 5 hours. Then, the precipitates were filtered with a Kiriyaama funnel under reduced pressure and washed with ether, and the thus obtained solid was dried at 40°C
15 for 10 hours under reduced pressure to obtain 5.20 g of N-methyl-N'-(1-naphthyl)urea (yield: 88%). Cyanoacetic acid (6.37 g, 74.9 mmol) and then acetic anhydride (7.65 g, 74.9 mmol) were added to a suspension of the obtained urea (5.00 g, 25.0 mmol) in ethyl acetate (200
20 mL), and the resulting mixture was heated under reflux for 4 hours. After the temperature was lowered to room temperature, the reaction mixture was concentrated under reduced pressure and then water (150 mL) was added to the residue to precipitate a solid. The
25 precipitated solid was filtered with a Kiriyaama funnel under reduced pressure and the thus obtained solid is added to water (150 mL). A 1N aqueous sodium hydroxide

solution was added thereto until the pH became 10, and the resulting mixture was vigorously stirred at room temperature for 1 hour. This mixture was filtered with a Kiriyaama funnel under reduced pressure and washed
5 with ether, and the thus obtained solid was dried at 40°C for 15 hours under reduced pressure to obtain 5.00 g of the title compound (yield: 75%).

Reference Example 2: 6-amino-3-methyl-1-(1-naphthyl)-5-nitrosouracil

10 The compound obtained in Reference Example 1 (1.00 g, 4.12 mmol) was added to an aqueous solution (2 mL) of sodium nitrite (426 mg, 6.18 mmol), followed by adding thereto concentrated hydrochloric acid (515 µL) under ice-cooling, and the resulting mixture was
15 stirred for 10 hours. The pH of the reaction mixture was adjusted to 7 with a 5% aqueous sodium hydrogencarbonate solution. The precipitates were filtered with a Kiriyaama funnel under reduced pressure and dried at 40°C for 15 hours under reduced pressure
20 to obtain 1.03 g of the title compound (yield 84%).

Reference Example 3: 5,6-diamino-3-methyl-1-(1-naphthyl)uracil

A suspension of the compound obtained in Reference Example 2 (1.03 g, 3.48 mmol) and 5%
25 palladium/carbon (100 mg) in methanol (10 mL) was stirred under a hydrogen atmosphere for 10 hours.

After the catalyst was filtered off, the solvent was distilled off under reduced pressure and diethyl ether (10 mL) was added to the residue. The precipitates were filtered with a Kiriya funnel under reduced pressure and dried at 40°C for 10 hours under reduced pressure to obtain 583 mg of the title compound (yield 59%).

Reference Example 4: 2,3,6-trimethyl-4-(pivaloyloxy)phenol

10 Pyridine (17.0 mL, 210 mmol) was added to a suspension of 2,3,6-trimethylhydroquinone (10.0 g, 65.7 mmol) in methylene chloride (70 mL), and then a solution of pivaloyl chloride (8.1 mL, 65.7 mmol) in methylene chloride (60 mL) was slowly dropped thereinto
15 under ice-cooling. After completion of the dropping, the temperature was raised to room temperature and the mixture was stirred for 20 hours, followed by adding thereto acetic acid (3.76 mL) and water (150 mL). The organic layer was washed with water and then a 10%
20 aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column chromatography to obtain 10.4 g of the title compound (yield: 67%).

25 Reference Example 5: 4-benzyloxy-2,3,5-trimethylphenol

After 60% sodium hydride (609 mg, 15.2 mmol)

was washed several times with n-hexane, it was suspended in dimethylformamide (20 mL) and a solution of the compound obtained in Reference Example 4 (3.00 g, 12.7 mmol) in dimethylformamide (5 mL) was added dropwise thereto under ice-cooling. After completion of the dropwise addition, the resulting mixture was stirred for 30 minutes, followed by adding thereto a solution of benzyl bromide (2.82 g, 16.5 mmol) in dimethylformamide (5 mL), and the resulting mixture was stirred at room temperature for 4 hours. The solvent was distilled off under reduced pressure and a 5% aqueous citric acid solution was added to the residue, followed by extraction with diisopropyl ether. The organic layer was washed with water and then a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column chromatography to obtain 2.64 g of 1-benzyloxy-4-(pivaloyloxy)-2,3,6-trimethylbenzene (yield: 64%). Then, to a solution of this compound (2.60 g, 7.96 mmol) in dimethylformamide (10 mL) was added a suspension of potassium hydroxide (1.79 g, 31.8 mmol) in methanol under ice-cooling, and the temperature was raised to room temperature, followed by stirring for 12 hours.

The solvent was distilled off under reduced pressure and a 5% aqueous citric acid solution was added to the residue, followed by extraction with

diisopropyl ether. The organic layer was washed successively with a 5% aqueous sodium hydrogencarbonate solution, water and a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was dissolved in methanol, and then water was added thereto to precipitate a solid. The solid was filtered with a Kiriya funnel under reduced pressure and washed with n-hexane, and the thus obtained solid was dried at 40°C for 12 hours under reduced pressure to obtain 1.68 g of the title compound as a white solid (yield: 87%).

Reference Example 6: (4-benzyloxy-2,3,5-trimethyl)phenoxyacetic acid

Under ice-cooling, 60% sodium hydride (73 mg, 1.82 mmol) washed several times with n-hexane was slowly added to a solution of the compound obtained in Reference Example 5 (400 mg, 1.65 mmol) in dimethylformamide (16 mL). The temperature was raised to room temperature and the mixture was stirred for 30 minutes, and then a solution of ethyl bromoacetate (202 μ L, 1.82 mmol) in dimethylformamide (2 mL) was added thereto, followed by stirring at room temperature for 3 hours. The solvent was distilled off under reduced pressure and a 5% aqueous citric acid solution was added to the residue, followed by extraction with diisopropyl ether. The organic layer was washed with

water and then a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column chromatography to obtain 495 mg of ethyl (4-benzyloxy-2,3,5-trimethyl)phenoxyacetate (yield: 91%). Then, to a solution of this ester (477 mg, 1.45 mmol) in ethanol (20 mL) was added a 1N aqueous sodium hydroxide solution (5 mL), and the resulting mixture was stirred at room temperature for 8 hours. After the pH was adjusted to 7 with 1N hydrochloric acid under ice-cooling, the solvent was distilled off under reduced pressure. Water was added to the residue to precipitate a solid and the solid was filtered with a Kiriyaama funnel under reduced pressure. The solid thus obtained was suspended in ethanol/water at room temperature for 1 hour and the suspension was filtered with a Kiriyaama funnel under reduced pressure. The precipitate was dried at 40°C for 15 hours under reduced pressure to obtain 415 mg of the title compound (yield: 95%).

Reference Example 7: tert-butyl 4-hydroxy-2,3,6-trimethylphenylcarbamate

Di-tert-butyl dicarbonate (1.51 g, 6.91 mmol) was added to a solution of 4-amino-2,3,5-trimethylphenol (950 mg, 6.28 mmol) in tetrahydrofuran (12 mL) and the resulting mixture was heated under

reflux for 2 hours. After the temperature was lowered to room temperature, water was added to the mixture, followed by extraction with diisopropyl ether. The organic layer was washed with water and then a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column chromatography to obtain 1.40 g of the title compound (yield: 89%).

10 Reference Example 8: tert-butyl 4-(carboxymethoxy)-2,3,6-trimethylphenylcarbamate

The title compound was obtained by the same process as in Reference Example 6 except for using the compound obtained in Reference Example 7, as a starting material in place of the compound obtained in Reference Example 5.

Reference Example 9: 6-amino-5-(6-benzyloxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-3-methyl-1-(1-naphthyl)uracil

20 Triethylamine (77 μ L, 0.55 mmol) and diphenylphosphoryl chloride (148 mg, 0.55 mmol) were added to a solution of 6-benzyloxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxylic acid (180 mg, 0.55 mmol) in ethyl acetate (6 mL) under ice-cooling, and the resulting mixture was stirred for 1 hour. Subsequently, the compound obtained in Reference

Example 3 (142 mg, 0.50 mmol) and then triethylamine (77 μ L, 0.55 mmol) were added thereto, followed by stirring at room temperature for 5 hours. The solvent was distilled off under reduced pressure and water was added to the residue, followed by extraction with methylene chloride. The organic layer was washed successively with a 5% aqueous sodium hydrogencarbonate solution, water and a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column chromatography to obtain 224 mg of the title compound (yield: 69%).

Reference Example 10: 6-amino-5-(6-benzyloxy-2,2,5,7-tetramethylchroman-8-carboxamide)-3-methyl-1-(1-naphthyl)uracil

The title compound was obtained by the same process as in Reference Example 9 except for using 6-benzyloxy-2,2,5,7-tetramethylchroman-8-carboxylic acid as a starting material in place of 6-benzyloxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxylic acid.

Reference Example 11: 6-amino-5-[(4-benzyloxy-2,3,5-trimethyl)phenoxy]acetamide-3-methyl-1-(1-naphthyl)uracil

To a solution of the compound obtained in Reference Example 3 (230 mg, 0.81 mmol) and the

compound obtained in Reference Example 6 (243 mg, 0.81 mmol) in dimethylformamide (6 mL) were added 1-hydroxybenzotriazole (120 mg, 0.89 mmol) and then N-ethyl-N'-[3-(dimethylamino)propyl]carbodiimide (171 mg, 0.89 mmol), and the resulting mixture was stirred at room temperature for 10 hours. The solvent was distilled off under reduced pressure and a 5% aqueous sodium hydrogencarbonate solution was added to the residue, followed by extraction with methylene chloride. The organic layer was washed successively with a 5% aqueous citric acid solution, water and a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column chromatography to obtain 242 mg of the title compound (yield: 53%).

Reference Example 12: 6-amino-5-[[4-(tert-butoxycarbonyl-amino)-2,3,5-trimethyl]phenoxy]acetamide-3-methyl-1-(1-naphthyl)uracil

The title compound was obtained by the same process as in Reference Example 15 except for using the compound obtained in Reference Example 8, as a starting material in place of the compound obtained in Reference Example 6.

Reference Example 13: 6-amino-5-(6-benzyloxy-2,2,5,7-

tetramethyl-3-oxaindan-4-yl)aminocarbonyl-3-methyl-1-
(1-naphthyl)uracil

Triethylamine (542 μ L, 3.89 mmol) and diphenylphosphoryl azide (1.07 g, 3.89 mmol) were added
5 to a solution of 6-benzyloxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxylic acid (1.00 g, 3.54 mmol) in toluene (20 mL) under ice-cooling, and the resulting mixture was stirred for 2 hours. Thereafter, the reaction temperature was raised to 90°C and the mixture
10 was stirred for another 3 hours and then cooled to room temperature to prepare a solution of an isocyanate corresponding to the starting carboxylic acid in toluene. On the other hand, 60% sodium hydride (113 mg, 2.83 mmol) washed several times with n-hexane was
15 slowly added to a solution of the compound obtained in Reference Example 1 (757 mg, 2.83 mmol) in dimethylformamide (20 mL) under ice-cooling. Subsequently, the above-mentioned solution of the isocyanate in toluene was added thereto and the
20 reaction was carried out at room temperature for 10 hours. The solvent was distilled off under reduced pressure and a 1N aqueous sodium hydroxide solution was added to the residue, followed by extraction with methylene chloride. The organic layer was washed with
25 water and then a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column

chromatography to obtain 250 mg of the title compound (yield: 15%).

Reference Example 14: 6-amino-5-(6-benzyloxy-2,2,5,7-tetramethyl-3-oxaindan-4-yl)carbonyl-3-methyl-1-(1-

5 naphthyl)uracil

A solution of oxalyl chloride (170 μ L, 1.95 mmol) in methylene (5 mL) chloride was added to a solution of 6-benzyloxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxylic acid (1.00 g, 3.54 mmol) in methylene
10 chloride (15 mL), and the resulting mixture was stirred under ice-cooling for 2 hours. The temperature was raised to room temperature to prepare a solution of an acid chloride corresponding to the starting carboxylic acid in methylene chloride. On the other hand, 60%
15 sodium hydride (170 mg, 4.25 mmol) washed several times with n-hexane was slowly added to a solution of the compound obtained in Reference Example 1 (1.14 g, 4.25 mmol) in dimethylformamide (15 mL) under ice-cooling. Subsequently, the above-mentioned solution of the acid
20 chloride in methylene chloride was added thereto and the reaction was carried out at room temperature for 10 hours. After the solvent was distilled off under reduced pressure, water was added to the residue and the precipitates were filtered under reduced pressure.
25 The thus obtained solid was purified by a silica gel column chromatography to obtain 650 mg of the title compound (yield: 32%).

Example 1: 6-amino-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-3-methyl-1-(1-naphthyl)uracil
(compound 1)

Concentrated hydrochloric acid (28 μ L) was
5 added to a suspension of the compound obtained in
Reference Example 9 (200 mg, 0.34 mmol) and 5%
palladium/carbon (20 mg) in methanol, and the resulting
mixture was stirred under a hydrogen atmosphere for 15
hours. After the catalyst was filtered off, the
10 solvent was distilled off under reduced pressure and
the residue was purified by a silica gel column
chromatography to obtain 140 mg of the title compound
(yield: 82%).

^1H NMR (DMSO- d_6) δ 8.76 (s, 1H), 8.18-8.10 (m, 2H),
15 7.75-7.55 (m, 6H), 5.91 (br s, 2H), 3.20 (s, 3H), 2.86
(s, 2H), 2.15 (s, 3H), 2.05 (s, 3H), 1.29 (s, 3H), 1.26
(s, 3H).

Example 2: 6-amino-5-(6-hydroxy-2,2,5,7-tetramethyl-
chroman-8-carboxamide)-3-methyl-1-(1-naphthyl)uracil

20 (compound 2)

The title compound was obtained by the same
process as in Example 1.

^1H NMR (CDCl₃) δ 8.08-7.97 (m, 2H), 7.77-7.75 (m, 1H),
7.65-7.60 (m, 4H), 7.24 (s, 1H), 4.99 (br s, 2H), 3.69
25 (s, 1H), 3.41 (s, 3H), 2.62 (t, J = 6.8 Hz, 2H), 2.29
(s, 3H), 2.14 (s, 3H), 1.78 (t, J = 6.8 Hz, 2H), 1.22

(s, 3H), 1.21 (s, 3H).

Example 3: 6-amino-5-(6-hydroxy-2,2,5,7-tetramethyl-chroman-8-carboxamide)-1-(1-naphthyl)uracil (compound 10)

5 The title compound was obtained by the same process as in Example 1.

¹H NMR (DMSO-d₆) δ 10.93 (s, 1H), 8.74 (s, 1H), 8.16-8.09 (m, 2H), 7.71-7.62 (m, 6H), 5.69 (br s, 2H), 2.51-2.50 (m, 2H), 2.14 (s, 3H), 2.06 (s, 3H), 1.70-1.67 (m, 2H),
10 1.09 (s, 3H), 1.08 (s, 3H).

Example 4: 6-amino-5-[(4-hydroxy-2,3,5-trimethyl)phenoxy]acetamide-3-methyl-1-(1-naphthyl)uracil (compound 3)

15 The title compound was obtained by the same process as in Example 1.

¹H NMR (DMSO-d₆) δ 8.44 (s, 1H), 8.14-8.08 (m, 2H), 7.70-7.54 (m, 6H), 6.53 (s, 1H), 6.13 (br s, 2H), 4.49 (s, 2H), 3.18 (s, 3H), 2.13 (s, 3H), 2.12 (s, 3H), 2.09 (s, 3H).

20 Example 5: 6-amino-5-[(4-amino-2,3,5-trimethyl)phenoxy]acetamide-3-methyl-1-(1-naphthyl)uracil (compound 4)

To a suspension of the compound obtained in Reference Example 12 (150 mg, 0.26 mmol) in ethyl
25 acetate (5 mL) were added 4N hydrochloric acid/1,4-

dioxane (1.2 mL) and acetic acid (3 mL), and the resulting mixture was stirred at room temperature for 2 hours. The solvent was distilled off under reduced pressure and a 5% aqueous sodium hydrogencarbonate solution was added to the residue, followed by extraction with methylene chloride. The organic layer was washed with water and then a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column chromatography to obtain 94 mg of the title compound (yield: 76%).

¹H NMR (DMSO-d₆) δ 8.41 (s, 1H), 8.14-8.08 (m, 2H), 7.70-7.56 (m, 5H), 6.61 (s, 1H), 6.12 (br s, 2H), 4.43 (s, 2H), 4.12 (br s, 2H), 3.18 (s, 3H), 2.12 (s, 3H), 2.06 (s, 3H), 2.01 (s, 3H).

Example 6: 6-amino-5-[[[(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-yl)methyl]amino]-3-methyl-1-(1-naphthyl)uracil (compound 15)

A borane-methyl sulfide complex (900 µL) was added to a solution of the compound obtained in Example 1 (950 mg, 1.90 mmol) in tetrahydrofuran (38 mL) under ice-cooling, and the resulting mixture was heated under reflux for 12 hours. Under ice-cooling, the reaction mixture was adjusted to pH 8 with a 1N aqueous sodium hydroxide solution and then extracted with methylene chloride. The organic layer was washed with water and

then a 10% aqueous sodium chloride solution and dried over anhydrous sodium sulfate. The solvent was distilled off under reduced pressure and the residue was purified by a silica gel column chromatography to
5 obtain 285 mg of the title compound (yield: 31%).

¹H NMR (DMSO-d₆) δ 8.12-8.06 (m, 2H), 7.68-7.42 (m, 7H), 5.74 (br s, 2H), 3.85-3.79 (m, 1H), 3.72-3.66 (m, 1H), 2.95 (s, 3H), 2.83 (s, 2H), 2.19 (s, 3H), 2.01 (s, 3H), 1.30 (s, 3H), 1.27 (s, 3H).

10 Example 7: 6-amino-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-yl)aminocarbonyl-3-methyl-1-(1-naphthyl)uracil (compound 16)

The title compound was obtained by the same process as in Example 1 except for using the compound
15 of Reference Example 13.

¹H NMR (DMSO-d₆) δ 8.50 (br s, 1H), 8.11-8.09 (m, 2H), 7.62-7.45 (m, 7H), 6.63 (s, 1H), 3.17 (s, 3H), 2.82 (s, 2H), 2.00 (s, 3H), 1.77 (s, 3H), 1.28 (s, 6H).

Example 8: 6-amino-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-yl)carbonyl-3-methyl-1-(1-naphthyl)uracil
20 (compound 17)

The title compound was obtained by the same process as in Example 1 except for using the compound of Reference Example 14.

25 ¹H NMR (DMSO-d₆) δ 8.67 (s, 1H), 8.11-8.06 (m, 2H), 7.83 (s, 1H), 7.73-7.62 (m, 5H), 6.65 (s, 1H), 3.20 (s, 3H),

2.65 (s, 2H), 1.95 (s, 6H), 0.87 (s, 3H), 0.75 (s, 3H).

Example 9: 6-amino-3-ethyl-5-(6-hydroxy-2,2,5,7-tetramethylchroman-8-carboxamide)-1-(6-methoxypyridin-3-yl)uracil (compound 439)

5 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆) δ 8.79 (s, 1H), 8.20 (d, J = 2.4 Hz, 1H), 7.73-7.78 (m, 2H), 6.99 (d, J = 8.9 Hz, 1H), 5.85 (br s, 2H), 3.94 (s, 3H), 3.83 (q, J = 6.9 Hz, 2H),
10 2.54 (m, 2H), 2.14 (s, 3H), 2.07 (s, 3H), 1.71 (t, J = 6.9 Hz, 3H), 1.07-1.15 (m, 8H).

Example 10: 6-amino-3-ethyl-5-(6-hydroxy-2,2,5,7-tetramethylchroman-8-carboxamide)-1-(2-methoxypyridin-3-yl)uracil (compound 433)

15 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆) δ 8.76 (s, 1H), 8.35 (m, 1H), 7.82 (q, J = 1.6 Hz, 1H), 7.73 (s, 1H), 7.16-7.21 (m, 1H), 5.84 (br s, 2H), 3.90 (s, 3H), 3.82 (q, J = 6.9 Hz, 2H),
20 2.54 (m, 2H), 2.14 (s, 3H), 2.06 (s, 3H), 1.72 (t, J = 6.9 Hz, 3H), 1.07-1.15 (m, 8H).

Example 11: 6-amino-1-(6-chloropyridin-3-yl)-3-ethyl-5-(6-hydroxy-2,2,5,7-tetramethylchroman-8-carboxamide)uracil (compound 457)

25 The title compound was obtained by the same

process as in Example 1.

1H NMR (DMSO-d₆) δ 8.84 (s, 1H), 8.52 (d, J = 2.4 Hz, 1H), 7.99-8.03 (m, 1H), 7.77 (s, 1H), 7.74 (s, 1H), 5.88 (br s, 2H), 3.82 (q, J = 6.9 Hz, 2H), 2.54 (m, 2H), 2.14 (s, 3H), 2.07 (s, 3H), 1.72 (t, J = 6.9 Hz, 3H), 1.07-1.14 (m, 8H).

Example 12: 6-amino-3-ethyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(1-naphthyl)uracil (compound 19)

10 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆) δ 1.13 (3H, t, J = 7.0 Hz), 1.25 (3H, s), 1.29 (3H, s), 2.05 (3H, s), 2.15 (3H, s), 2.85 (2H, s), 3.86 (2H, q, J = 7.0 Hz), 5.91 (2H, brs), 7.53-7.55
15 (1H, m), 7.62-7.72 (4H, m), 7.76 (1H, s), 8.10-8.17 (2H, m), 8.74 (1H, s); MS m/z 515(M+H)⁺

Example 13: 6-amino-3-propyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(1-naphthyl)uracil (compound 20)

20 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆) δ 0.86 (3H, t, J = 7.4 Hz), 1.25 (3H, s), 1.29 (3H, s), 1.54-1.60 (2H, m), 2.05 (3H, s), 2.15 (3H, s), 2.85 (2H, s), 3.78 (2H, t, J = 7.4 Hz), 5.92
25 (2H, brs), 7.52-7.54 (1H, m), 7.62-7.72 (4H, m), 7.75 (1H, s), 8.10-8.17 (2H, m), 8.74 (1H, s); MS m/z

529 (M+H)⁺

Example 14: 6-amino-3-butyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(1-naphthyl)uracil (compound 21)

5 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆) δ 0.89 (3H, t, J = 7.3 Hz), 1.25-1.34 (8H, m), 1.50-1.57 (2H, m), 2.05 (3H, s), 2.15 (3H, s), 2.85 (2H, s), 3.81 (2H, t, J = 7.3 Hz), 5.92 (2H, brs),
10 7.51-7.54 (1H, m), 7.62-7.72 (4H, m), 7.75 (1H, s), 8.10-8.17 (2H, m), 8.74 (1H, s); MS m/z 543 (M+H)⁺

Example 15: 6-methylamino-3-methyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(1-naphthyl)uracil (compound 502)

15 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆) δ 1.27 (6H, s), 2.06 (3H, s), 2.31 (3H, s), 2.82 (3H, d, J = 5.1 Hz), 2.84 (2H, s), 3.18 (3H, s), 5.29-5.33 (1H, m), 7.55-7.68 (6H, m), 8.08-8.14
20 (2H, m), 8.75 (1H, s); MS m/z 515 (M+H)⁺

Example 16: 6-amino-3-methyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(2,3-dimethylphenyl)uracil (compound 301)

25 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d6) δ 1.34 (6H, s), 1.99 (3H, s), 2.06 (3H, s), 2.13 (3H, s), 2.34 (3H, s), 2.88 (2H, s), 3.17 (3H, s), 5.83 (2H, brs), 7.12 (1H, d, J = 7.5 Hz), 7.26-7.30 (1H, m), 7.35 (1H, d, J = 7.4 Hz), 7.75 (1H, brs), 8.71
5 (1H, s); MS m/z 479(M+H)⁺

Example 17: 6-amino-3-ethyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(2,3-dimethyl-phenyl)uracil (compound 331)

The title compound was obtained by the same
10 process as in Example 1.

1H NMR (DMSO-d6) δ 1.10 (3H, t, J = 7.0 Hz), 1.34 (6H, s), 1.98 (3H, s), 2.06 (3H, s), 2.13 (3H, s), 2.34 (3H, s), 2.88 (2H, s), 3.85 (2H, q, J = 7.0 Hz), 5.83 (2H, brs), 7.13 (1H, d, J = 7.5 Hz), 7.26-7.30 (1H, m), 7.35
15 (1H, d, J = 7.4 Hz), 7.74 (1H, s), 8.68 (1H, s); MS m/z 493(M+H)⁺

Example 18: 6-amino-3-propyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(2,3-dimethyl-phenyl)uracil (compound 361)

20 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d6) δ 0.85 (3H, t, J = 7.4 Hz), 1.33 (3H, s), 1.34 (3H, s), 1.50-1.60 (2H, m), 1.98 (3H, s), 2.05 (3H, s), 2.13 (3H, s), 2.34 (3H, s), 2.87 (2H, s),
25 3.71-3.79 (2H, m), 5.83 (2H, brs), 7.13 (1H, d, J = 7.5 Hz), 7.26-7.30 (1H, m), 7.35 (1H, d, J = 7.4 Hz), 7.74

(1H, s), 8.68 (1H, s); MS m/z 507 (M+H)⁺

Example 19: 6-methylamino-3-methyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(2,3-dimethylphenyl)uracil (compound 506)

5 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆) δ 1.30 (6H, s), 1.96 (3H, s), 2.05 (3H, s), 2.28 (3H, s), 2.32 (3H, s), 2.85 (2H, s), 2.89 (3H, d, J = 5.1 Hz), 3.15 (3H, s), 5.07-5.10 (1H, m), 7.05
10 (1H, d, J = 7.5 Hz), 7.23-7.27 (1H, m), 7.32 (1H, d, J = 7.4 Hz), 7.65 (1H, s), 8.69 (1H, s); MS m/z 493 (M+H)⁺

Example 20: 6-amino-3-ethyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(4-indanyl)uracil (compound 91)

15 The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆) δ 1.10 (3H, t, J = 7.0 Hz), 1.34 (6H, s), 2.01-2.09 (5H, m), 2.13 (3H, s), 2.61-2.74 (2H, m), 2.88 (2H, s), 2.93-3.04 (2H, m), 3.83 (2H, q, J = 7.0
20 Hz), 5.88 (2H, brs), 7.12 (1H, d, J = 7.4 Hz), 7.31-7.35 (1H, m), 7.41 (1H, d, J = 7.3 Hz), 7.75 (1H, s), 8.67 (1H, s); MS m/z 505 (M+H)⁺

Example 21: 6-amino-3-propyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(4-indanyl)uracil (compound 121)

25

The title compound was obtained by the same process as in Example 1.

¹H NMR (DMSO-d₆) δ 0.85 (3H, t, J = 7.4 Hz), 1.34 (6H, s), 1.50-1.59 (2H, m), 2.01-2.09 (5H, m), 2.13 (3H, s), 2.60-2.74 (2H, m), 2.88 (2H, s), 2.93-3.05 (2H, m), 3.69-3.80 (2H, m), 5.88 (2H, brs), 7.11 (1H, d, J = 7.4 Hz), 7.31-7.35 (1H, m), 7.41 (1H, d, J = 7.4 Hz), 7.74 (1H, s), 8.67 (1H, s); MS m/z 519(M+H)⁺

Example 22: 6-amino-3-methyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(5,6,7,8-tetrahydro-1-naphthyl)uracil (compound 181)

The title compound was obtained by the same process as in Example 1.

¹H NMR (DMSO-d₆) δ 1.34 (3H, s), 1.35 (3H, s), 1.70-1.77 (4H, m), 2.06 (3H, s), 2.13 (3H, s), 2.40-2.41 (2H, m), 2.81-2.82 (2H, m), 2.88 (2H, s), 3.16 (3H, s), 5.82 (2H, brs), 7.09-7.11 (1H, m), 7.25-7.31 (2H, m), 7.74 (1H, s), 8.70 (1H, s); MS m/z 505(M+H)⁺

Example 23: 6-amino-3-ethyl-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(5,6,7,8-tetrahydro-1-naphthyl)uracil (compound 211)

The title compound was obtained by the same process as in Example 1.

¹H NMR (DMSO-d₆) δ 0.86 (3H, t, J = 6.8 Hz), 1.34 (6H, s), 1.71-1.74 (4H, m), 2.06 (3H, s), 2.13 (3H, s), 2.33-2.45 (2H, m), 2.76-2.82 (2H, m), 2.88 (2H, s),

5.82 (2H, brs), 7.09-7.12 (1H, m), 7.25-7.31 (2H, m),
7.74 (1H, s), 8.67 (1H, s); MS m/z 519(M+H)⁺

Example 24: 6-amino-3-propyl-5-(6-hydroxy-2,2,5,7-
tetramethyl-3-oxaindan-4-carboxamide)-1-(5,6,7,8-
5 tetrahydro-1-naphthyl)uracil (compound 241)

The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d₆)δ 0.85 (3H, t, J = 7.4 Hz), 1.34 (6H, s), 1.50-1.59 (2H, m), 1.70-1.77 (4H, m), 2.05 (3H, s),
10 2.12 (3H, s), 2.28-2.45 (2H, m), 2.76-2.82 (2H, m),
2.88 (2H, s), 5.83 (2H, brs), 7.09-7.13 (1H, m), 7.21-
7.31 (2H, m), 7.74 (1H, s), 8.67 (1H, s); MS m/z
533(M+H)⁺

Example 25: 6-amino-5-[(4-amino-2,3,5-
15 trimethyl)phenoxy]acetamide-3-ethyl-1-(1-
naphthyl)uracil (compound 12)

The title compound was obtained by the same process as in Example 5.

1H NMR (DMSO-d₆)δ 1.11 (3H, t, J = 6.9 Hz), 2.01 (3H, s),
20 s), 2.06 (3H, s), 2.13 (3H, s), 3.81-3.88 (2H, m), 4.12
(2H, brs), 4.42 (2H, s), 6.12 (2H, brs), 6.61 (1H, s),
7.52-7.69 (5H, m), 8.08-8.13 (2H, m), 8.42 (1H, s); MS
m/z 488(M+H)⁺

Example 26: 6-amino-5-[(4-amino-2,3,5-
25 trimethyl)phenoxy]acetamide-3-propyl-1-(1-

naphthyl)uracil (compound 13)

The title compound was obtained by the same process as in Example 5.

1H NMR (DMSO-d₆) δ 0.85 (3H, t, J = 7.5 Hz), 1.51-1.60
 5 (2H, m), 2.01 (3H, s), 2.06 (3H, s), 2.13 (3H, s), 3.76
 (2H, t, J = 7.3 Hz), 4.12 (2H, brs), 4.42 (2H, s), 6.12
 (2H, brs), 6.60 (1H, s), 7.51-7.69 (5H, m), 8.08-8.13
 (2H, m), 8.42 (1H, s); MS m/z 502 (M+H)⁺

10 Example 27: 6-amino-5-[(4-amino-2,3,5-
trimethyl)phenoxy]acetamide-3-butyl-1-(1-
naphthyl)uracil (compound 14)

The title compound was obtained by the same process as in Example 5.

15 1H NMR (DMSO-d₆) δ 0.88 (3H, t, J = 7.3 Hz), 1.23-1.32
 (2H, m), 1.48-1.55 (2H, m), 2.01 (3H, s), 2.06 (3H, s),
 2.13 (3H, s), 3.79 (2H, t, J = 7.3 Hz), 4.20 (2H, brs),
 4.42 (2H, s), 6.12 (2H, brs), 6.60 (1H, s), 7.52-7.69
 (5H, m), 8.08-8.13 (2H, m), 8.42 (1H, s); MS m/z 516
 20 (M+H)⁺

Example 28: 6-methylamino-5-[(4-amino-2,3,5-
trimethyl)phenoxy]acetamide-3-methyl-1-(1-
naphthyl)uracil (compound 487)

The title compound was obtained by the same
 25 process as in Example 5.

1H NMR (DMSO-d₆) δ 2.00 (3H, s), 2.04 (3H, s), 2.10 (3H,
 s), 2.60 (3H, d, J = 5.0 Hz), 4.13 (2H, brs), 4.42 (2H,

s), 5.40-5.44 (2H, m), 6.60 (1H, s), 7.54-7.68 (5H, m), 8.07-8.13 (2H, m), 8.71 (1H, s); MS m/z 488 (M+H)⁺

Example 29: 6-amino-5-[(4-amino-2,3,5-trimethyl)phenoxy]acetamide-3-methyl-1-(2,3-dimethylphenyl)uracil (compound 304)

The title compound was obtained by the same process as in Example 5.

¹H NMR (DMSO-d₆) δ 1.97 (3H, s), 2.01 (3H, s), 2.06 (3H, s), 2.12 (3H, s), 2.32 (3H, s), 3.15 (2H, s), 4.12 (2H, brs), 4.42 (2H, s), 6.00 (2H, brs), 6.60 (1H, s), 7.08 (1H, d, J = 7.6 Hz), 7.24-7.27 (1H, m), 7.32 (1H, d, J = 7.5 Hz), 8.38 (1H, s); MS m/z 452 (M+H)⁺

Example 30: 6-methylamino-5-[(4-amino-2,3,5-trimethyl)phenoxy]acetamide-3-methyl-1-(2,3-dimethylphenyl)uracil (compound 481)

The title compound was obtained by the same process as in Example 5.

¹H NMR (DMSO-d₆) δ 1.95 (3H, s), 2.00 (3H, s), 2.04 (3H, s), 2.10 (3H, s), 2.31 (3H, s), 2.67 (3H, t, J = 5.0 Hz), 3.14 (3H, s), 4.13 (2H, brs), 4.40 (2H, s), 5.20-5.24 (1H, m), 6.59 (1H, s), 7.05 (1H, d, J = 7.6 Hz), 7.22-7.26 (1H, m), 7.32 (1H, d, J = 7.5 Hz), 8.66 (1H, s); MS m/z 466 (M+H)⁺

Example 31: 6-ethylamino-5-[(4-amino-2,3,5-trimethyl)phenoxy]acetamide-3-methyl-1-(2,3-dimethylphenyl)uracil

(compound 482)

The title compound was obtained by the same process as in Example 5.

¹H NMR (DMSO-d₆) δ 0.84 (3H, t, J = 7.1 Hz), 1.96 (3H, s), 2.00 (3H, s), 2.05 (3H, s), 2.10 (3H, s), 2.31 (3H, s), 3.04-3.12 (2H, m), 3.15 (3H, s), 4.13 (2H, brs), 4.92-4.95 (1H, m), 6.59 (1H, s), 7.07 (1H, d, J = 7.4 Hz), 7.23-7.27 (1H, m), 7.32 (1H, d, J = 7.4 Hz), 8.68 (1H, s); MS m/z 480 (M+H)⁺

10 Example 32: 6-propylamino-5-[(4-amino-2,3,5-trimethyl)phenoxy]acetamide-3-methyl-1-(2,3-dimethylphenyl)uracil (compound 483)

The title compound was obtained by the same process as in Example 5.

15 ¹H NMR (DMSO-d₆) δ 0.59 (3H, t, J = 7.4 Hz), 1.22-1.29 (2H, m), 1.96 (3H, s), 2.00 (3H, s), 2.05 (3H, s), 2.10 (3H, s), 2.31 (3H, s), 2.95-3.01 (2H, m), 3.15 (3H, s), 4.13 (2H, brs), 4.40 (2H, s), 4.86-4.89 (1H, m), 6.60 (1H, s), 7.08 (1H, d, J = 7.6 Hz), 7.24-7.28 (1H, m),
20 7.33 (1H, d, J = 7.4 Hz), 8.67 (1H, s); MS m/z 494 (M+H)⁺

Example 33: 6-butylamino-5-[(4-amino-2,3,5-trimethyl)-phenoxy]acetamide-3-methyl-1-(2,3-dimethylphenyl)uracil
(compound 484)

25 The title compound was obtained by the same process as in Example 5.

1H NMR (DMSO-d6) δ 0.72 (3H, t, J = 7.3 Hz), 0.97-1.02 (2H, m), 1.19-1.24 (2H, m), 1.96 (3H, s), 2.00 (3H, s), 2.05 (3H, s), 2.10 (3H, s), 2.31 (3H, s), 2.98-3.04 (2H, m), 3.15 (3H, s), 4.14 (2H, brs), 4.40 (2H, s), 5 4.82-4.85 (1H, m), 6.60 (1H, s), 7.07 (1H, d, J = 7.5 Hz), 7.24-7.28 (1H, m), 7.33 (1H, d, J = 7.4 Hz), 8.66 (1H, s); MS m/z 508 (M+H)⁺

Example 34: 6-amino-3-(2-hydroxyethyl)-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(2,3-
10 dimethylphenyl)uracil (compound 466)

The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d6) δ 1.34 (6H, s), 1.98 (3H, s), 2.05 (3H, s), 2.13 (3H, s), 2.33 (3H, s), 2.87 (2H, s), 3.43-3.51
15 (2H, m), 3.88 (2H, t, J = 6.8 Hz), 4.75 (1H, t, J = 5.6 Hz), 5.83 (2H, brs), 7.11 (1H, d, J = 7.2 Hz), 7.28 (1H, t, J = 8.0 Hz), 7.35 (1H, d, J = 7.2 Hz), 7.74 (1H, s), 8.69 (1H, s); MS m/z 509 (M+H)⁺

Example 35: 6-amino-3-(3-hydroxypropyl)-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(2,3-
20 dimethylphenyl)uracil (compound 467)

The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d6) δ 1.34 (6H, s), 1.65-1.72 (2H, m), 1.98
25 (3H, s), 2.05 (3H, s), 2.13 (3H, s), 2.34 (3H, s), 2.88 (2H, s), 3.38-3.43 (2H, m), 3.80-3.88 (2H, m), 4.43

(1H, t, J = 5.4 Hz), 5.91 (2H, brs), 7.12 (1H, d, J = 6.0 Hz), 7.28 (1H, t, J = 7.6 Hz), 7.42 (1H, d, J = 6.0 Hz), 7.74 (1H, s), 8.73 (1H, s); MS m/z 523 (M+H)+

5 Example 36: 6-amino-3-(2-methoxyethyl)-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-carboxamide)-1-(2,3-dimethylphenyl)uracil (compound 464)

The title compound was obtained by the same process as in Example 1.

1H NMR (DMSO-d6) δ 1.34 (6H, s), 1.65-1.72 (2H, m), 1.98
10 (3H, s), 2.05 (3H, s), 2.13 (3H, s), 2.34 (3H, s), 2.88 (2H, s), 3.27 (3H, s), 3.48 (2H, t, J = 6.0 Hz), 3.95-3.99 (2H, m), 5.87 (2H, brs), 7.11 (1H, d, J = 7.6 Hz), 7.28 (1H, t, J = 7.6 Hz), 7.35 (1H, d, J = 7.6 Hz), 7.74 (1H, s), 8.69 (1H, s); MS m/z 523 (M+H)+

15 Example 37: 6-amino-1-(2,3-dimethylphenyl)-5-(6-hydroxy-2,2,5,7-tetramethyl-3-oxaindan-4-yl)aminocarbonyl-3-(2,2,2-trifluoroethyl)uracil

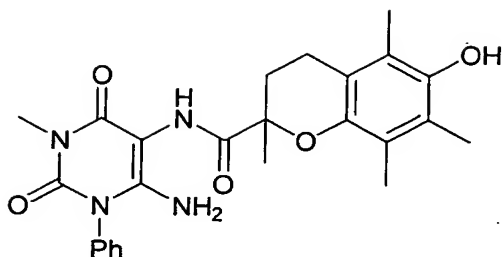
The title compound was obtained by the same process as in Example 1.

20 1H NMR (DMSO-d6) δ 1.34 (6H, s), 1.99 (3H, s), 2.06 (3H, s), 2.12 (3H, s), 2.35 (3H, s), 2.88 (2H, s), 4.57-4.69 (2H, m), 6.10 (2H, brs), 7.15 (1H, d, J = 7.6 Hz), 7.28-7.32 (1H, m), 7.38 (1H, d, J = 7.5 Hz), 7.76 (1H, s), 8.77 (1H, s)

25 Evaluation Example 1: Inhibitory effect on dermatitis

induced by picryl chloride

In order to verify the inflammation-inhibiting effect of the uracil derivative of the present invention, its effect on dermatitis induced by picryl chloride, a model of typical type IV allergic inflammation was evaluated by adopting the method of Asherson et al. (Immunology, 15, 405 (1968)). On the abdominal skin of each of ICR strain male mice was applied 0.1 ml of a 7% (w/v) picryl chloride/acetone solution to induce sensitization. On the seventh days after the sensitization, 0.02 ml of a 1% (w/v) picryl chloride/acetone solution was applied on the ear of each mouse to cause an allergic reaction. Immediately after this causing, 0.04 ml of acetone (a control) or a 0.25% (w/v) test compound/acetone solution was applied on the ear. Twenty-four hours after the causing, the thickness of the ear was measured and the dermatitis-inhibiting effect of the test compound was evaluated by employing the difference between the thickness of the ear 24 hours after the causing and that before the causing as an indication. The thickness of the ears of the mice treated with the test compound was compared with the thickness of the ears of mice on which 0.04 ml of a 2.5% (w/v) solution of the comparative compound 1 (the following structure) disclosed in Japanese Patent No. 3093170, as a comparative example, in acetone had been applied:



Comparative compound 1

From the difference of the ear thickness before and after the causing of the allergic reaction in the case of each group, the relative degree of efficacy of the test compound in the inhibition of ear enlargement was calculated according to the following equation by comparison with a group treated with the comparative compound in an amount of 10 times that of the test compound:

$$\begin{array}{l}
 \text{Degree of efficacy} = \frac{\left(\begin{array}{l} \text{Difference of ear thickness in the} \\ \text{case of control group} \end{array} \right) - \left(\begin{array}{l} \text{Difference of ear thickness in the} \\ \text{case of group treated with test compound} \end{array} \right)}{\left(\begin{array}{l} \text{Difference of ear thickness in the} \\ \text{case of control group} \end{array} \right) - \left(\begin{array}{l} \text{Difference of ear thickness in the} \\ \text{case of comparative group} \end{array} \right)}
 \end{array}$$

As a result, it was found that as shown in Table 19, the degree of efficacy of the uracil derivatives of the present invention was about 1. This fact indicates that the uracil derivatives of the present invention exhibit enlargement-inhibiting effect substantially equal to that obtained by the application

of the compound as comparative example in an amount of 10 times that of each uracil derivative of the present invention. Therefore, it was revealed that the uracil derivatives of the present invention are superior to
5 the compound as comparative example in anti-inflammatory effect on type IV allergic inflammation.

Table 19

Test compound	Degree of efficacy
Example 4	0.9
Example 5	1.1

Evaluation Example 2: Inhibitory effect on dermatitis induced by picryl chloride

10 The effect on the animal model in Evaluation Example 1 was verified by oral administration. ICR strain male mice were sensitized in the same manner as in Evaluation Example 1, and on the seventh days after the sensitization, 0.02 ml of a 1% (w/v) picryl
15 chloride/acetone solution was applied on the ear of each mouse to cause an allergic reaction. Each test compound was suspended in a 0.5% CMC-Na solution and orally administered in a dose of 10 mg/kg one hour before the causing of the allergic reaction. As a
20 comparative example, the above-mentioned comparative compound 1 disclosed in Japanese Patent No. 3093170 was orally administered in a dose of 100 mg/kg to make a comparison.

As a result, it was found that as shown in Table 20, the degree of efficacy of the uracil derivatives of the present invention was 1.0 to 1.6. This fact indicates that the uracil derivatives of the present invention exhibit enlargement-inhibiting effect substantially equal to that obtained by the administration of the compound as comparative example in an amount of 10 times that of each uracil derivative of the present invention. Therefore, it was revealed that the uracil derivatives of the present invention are superior to the compound as comparative example in anti-inflammatory effect on type IV allergic inflammation.

Table 20

Test compound	Degree of efficacy
Example 10	1.6
Example 14	1.0
Example 20	1.2
Example 22	1.3
Example 29	1.4
Example 30	1.6

Evaluation Example 3: Inhibitory effect on the itch-related behavior of mice induced by substance P

It has been reported that in the lesion part of a patient with atopic dermatitis, there are observed an increase in nerve fiber containing substance P (SP), a nerve peptide (Tobin D et al., J Allergy Clin

immunol, 90, 613-22(1992)) and an increase in reactivity with SP (Gianetti A et al. Br J Dermatol, 121, 681-8(1989)). On the other hand, it has been reported that when SP is administered to the

5 cervicodorsal part of a mouse, itch-related behavior is induced (Kuraishi Y et al. Eur J Pharmacol, 275, 229-33(1995)) and is suppressed by a certain antiallergic agent (Inagaki N et al. Eur J Pharmacol, 400, 73-9(2000)). Therefore, in order to verify the itch-

10 inhibiting effect of the uracil derivatives of the present invention, the itch-inhibiting effect was evaluated by employing the effect of the uracil derivatives on itch-related behavior induced by SP, as an indication. Each of BALB/c strain male mice was

15 previously transferred to a cage for observation and allowed to acclimate to an observation environment, and then SP was subcutaneously administered (200 µg/mouse) to the cervicodorsal part of the mouse. After the administration, the mouse was returned to the cage for

20 observation and the frequency of its itch-related behavior was measured for 60 minutes after the return. Each test drug was suspended in a 0.5% CMC-Na solution and orally administered in a dose of 10 mg/kg 30 minutes before the administration of SP. For

25 evaluating the itch-inhibiting effect, the inhibition rate of the test drug was calculated as an indication as follows by comparison with a group treated with a 0.5% CMC-Na solution. As a comparative example, an

antiallergic agent Oxatomide, which is considered as an agent having itch-inhibiting effect, was administered in a dose of 3 times that of the test drug (30 mg/kg).

$$\text{Inhibition rate} = \frac{\left(\begin{array}{l} \text{Frequency in the} \\ \text{case of group} \\ \text{treated with 0.5\%} \\ \text{CMC-Na solution} \end{array} \right) - \left(\begin{array}{l} \text{Frequency in the} \\ \text{case of group} \\ \text{treated with test} \\ \text{drug} \end{array} \right)}{\left(\begin{array}{l} \text{Frequency in the} \\ \text{case of group} \\ \text{treated with 0.5\%} \\ \text{CMC-Na solution} \end{array} \right)} \times 100$$

As a result, it was found that as shown in Table 21, the degree of the itch-inhibiting effect of the uracil derivatives of the present invention is equal to or larger than that of the antiallergic agent Oxatomide.

Table 21

Test compound	Inhibition rate (%)
Oxatomide 30 mg/kg	39
Example 2 10 mg/kg	71
Example 5 10 mg/kg	53

10 Formulation Example 1: Water-soluble ointment

A water-soluble ointment having the following composition was prepared by a conventional process:

Ingredients	Content per 2 g of ointment
Compound of Example 4	40 mg

Poly(ethylene glycol) 400	1372 mg
Poly(ethylene glycol) 4000	588 mg

Formulation Example 2: Tablets for internal use

Tablets for internal use having the following
5 composition was prepared by a conventional process:

Ingredients	Amount (mg/tablet)
Compound of Example 5	100
Lactose	353
Carboxymethyl cellulose calcium	30
10 Hydroxypropyl cellulose	7
Magnesium stearate	5
Crystalline cellulose	5
Total	500 mg

INDUSTRIAL APPLICABILITY

15 The uracil derivatives of the present
invention have not only a marked inflammation-
inhibiting effect on allergic inflammation, in
particular, type IV allergic inflammation, but also
itch-inhibiting effect. The uracil derivatives of the
20 present invention suppress type IV allergic
inflammation effectively and hence are useful as a
therapeutic agent for allergic diseases, in particular,
type IV allergic diseases. They can also suppress itch
and hence are useful also as anti-itch agent. They are
25 very useful particularly as a therapeutic agent for
atopic dermatitis from the viewpoint of both anti-

inflammation effect and anti-itch effect. In addition, since they are absorbed through skin when administered as an external preparation, they are useful for treating skin diseases such as atopic dermatitis, 5 contact dermatitis, psoriasis and the like. Furthermore, said uracil derivatives are advantageous in that they are nonsteroidal substances and hence do not exhibit any adverse effect similar to that of steroids.